







Db 48613 AATAAGGCATAGACTTCAACCTTTTAATAGTAAATAAAATTAAGTGAAAC 48672  
 Qy 1399 TGCACCTTAATGTTAAAGAGTTAGTGTGCTTAACTATGCAACCTGAAATA 1458  
 Db 48673 TGCACCTTGTAAATGTTAAAGAGTTAGTGTGCTTAACTATGCAACCTGAAAT 48732  
 Qy 1459 .GCTGTAAATGTCACAGAGATAATTCTAGCTTGTAGCTTAAGAATTGAGCAGGTGTT 1518  
 Db 48733 GCTGTAAATGTCACAGAGATAATTCTAGCTTGTAGCTTAAGAATTGAGCAGGTGTT 48792  
 Qy 1519 GTTGGGAGACTGCTGAGTCAACCCAAATAGTTGTTGAGGACTGGAGATGTTG 1578  
 Db 48793 GTTGGGAGACTGCTGAGTCAACCCAAATAGTTGTTGAGGACTGGAGATGTTG 48852  
 Qy 1579 ATCTGGGGCACATAGCCATTGCTGAGTCAAGCTTGTGTTGAAATCA 1638  
 Db 48853 ATCTGGGGCACATTGCTGAGTCAAGCTTGTGAGGCAATTGATGTTGAAATCA 48912  
 Qy 1639 CAGTATACGCCATGGCTCATCTCAGCTGGATCCTCCAGGCTTGCCTGCCA 1698  
 Db 48913 CAGTATACGCCATGGCTCATCTCAGCTGGATCCTCCAGGCTTGCCTGCCA 48972  
 Qy 1699 AAAGCCTTTGTGTTGTATCATATGAAGCTCATGGTTAAATCACATTGAGT 1758  
 Db 48973 AAAGCCTTTGTGTTGTATCATATGAAGCTCATGGTTAAATCACATTGAGT 49032  
 Qy 1759 GTTTCAGTGGTCCGAGATGCTGTGCTGATGCTCATATGTTGCAAGTGGAA 1818  
 Db 49033 GTTTCAGTGGTCCGAGATGCTGTGCTGATGCTCATATGTTGCAAGTGGAA 49092  
 Qy 1819 CTCCCTAAATAAAATGGCTCTAACTTAAACCCATTGTAAGAAATGGAAAG 1878  
 Db 49093 CTCCCTAAATAAAATGGCTCTAACTTAAACCCATTGTAAGAAATGGAAAG 49152  
 Qy 1879 GTGGAAAGCTCCCTGAAGTAAGCAAGACTTCTCTTACTGAGCCAACTTAAAGATGGAAG 1938  
 Db 49153 GTGGAAAGCTCCCTGAAGAAAGACTTCTCTTACTGAGCCAACTTAAAGATGGAAG 49212  
 Qy 1939 TCTCTATGTCGCCAGTGTGTTCTGATCGATGAAAGCAACCTGCTCTAG 1998  
 Db 49213 TCTCTATGTCGCCAGTGTGTTCTGATCGATGAAAGCAACCTGCTCTAG 49272  
 Qy 1999 ACCAGGCAACTTGGAAACTTGAAGCTCCAAACTGGACTATGGCTTACCCACAT 2058  
 Db 49273 ACCAGGCAACTTGGAAACTTGAAGCTCCAAACTGGACTATGGCTTACCCACAT 49332  
 Qy 2059 GGCTAAAGGGTTCAAGAAAGTGGGACAGGAACTTCACTTCATATATT 2118  
 Db 49333 GGCTAAAGGGTTCAAGAAAGTGGGACAGGAACTTCACTTCATATATT 49392  
 Qy 2119 GTATGATCTTATGAATGCTAAATGTTAAGTGTGTTGATGAAATTAATCTGT 2178  
 Db 49393 GTATGATCTTATGAATGCTAAATGTTAAGTGTGTTGATGAAATTAATCTGT 49452  
 Qy 2179 TTTAACAACTATGATTGTTGAAATAATCAATGCTTAACTATGTTGAAAG 2232  
 Db 49453 TTTAACAACTATGATTGTTGAAATAATCAATGCTTAACTATGTTGAAAG 49506

RESULT 3  
 ID AAF21272 standard; DNA; 143068 BP.  
 XX  
 AC AAF21272;  
 XX  
 DT 14-MAR-2001 (first entry)  
 DE Human low adenosine antisense oligonucleotide related sequence #2839.  
 XX  
 KW Low adenosine antisense oligonucleotide; phosphorothioate; allergy;  
 XX  
 KW airway disorder; bronchoconstriction; lung inflammation;  
 KW surfactant depletion; respiratory; bronchodilator; antiinflammatory;

KW immuno suppressive; antiasthmatic; analgesic; hypotensive; cytostatic;  
 KW respiratory obstruction; pulmonary obstruction; impeded respiration;  
 KW respiratory hypoproduction; Pulmonary vasoconstriction; asthma; RDS;  
 KW respiratory distress syndrome; pain; cystic fibrosis; allergic rhinitis;  
 KW pulmonary hypertension; emphysema; pulmonary transplantation rejection;  
 KW chronic obstructive pulmonary disease; pulmonary infection; bronchitis;  
 KW cancer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO20062736-A2.  
 XX  
 PD 26-OCT-2000.  
 XX  
 PF 24-MAR-2000; 2000WO-US08020.  
 XX  
 PR 06-APR-1999; 99US-0127958.  
 XX  
 (UYEC-) UNIV EAST CAROLINA.  
 PA (NYCE-) NYCE J W.  
 XX  
 PI NYCE JW;  
 XX  
 DR 2000-679539/66.  
 XX  
 PT Low adenosine (A) content antisense oligonucleotides which do not  
 trigger adenosine receptors during metabolism, useful e.g. for treating  
 cancers and respiratory obstructions -  
 XX  
 Disclosure; Page 1186-1219; 1592pp; English.  
 XX  
 PS Disclosure; Page 1186-1219; 1592pp; English.  
 XX  
 CC The present invention describes low adenosine (A) content antisense  
 oligonucleotides and compositions (I) comprising them. In the antisense  
 oligonucleotides the A is replaced by a 'universal', or alternative base.  
 CC (I) can have respiratory, bronchiodilator, antifibrillatory, analgesic,  
 CC immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.  
 CC The antisense oligonucleotides and (I) can be used to down-regulate the  
 CC expression and/or activity of target polypeptides associated with the  
 CC lung/respiratory disorders and malignancies, such as stimulating and  
 CC activating peptide factors and transmitters, transcription factors,  
 CC immunoglobulins and antibodies, antibody receptors, cytokines and  
 CC chemokines, endogenously produced specific and non-specific enzymes,  
 CC binding proteins, adhesion molecules and their receptors, cytokine and  
 CC chemokine receptors, adenosine receptors, bradykinin receptors, central  
 CC nervous system (CNS) and peripheral nervous and non-nervous system  
 CC transmitters, CNS and peripheral nervous and non-nervous system peptide  
 CC receptors, growth factors, vasoactive peptides and  
 CC receptors, binding proteins and malignancy associated proteins. The  
 CC antisense oligonucleotides may be used in this way to treat disorders  
 CC including respiratory obstruction (especially pulmonary obstruction  
 CC and/or bronchoconstriction) and/or lung inflammation, allergy(ies)  
 CC and/or surfactant hypoproduction which are associated with a disease or  
 CC condition selected from pulmonary vasoconstriction, inflammation, or  
 CC allergies, asthma, impeded respiration, respiratory distress syndrome  
 CC (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary  
 CC hypertension, emphysema, chronic obstructive pulmonary disease (COPD),  
 CC pulmonary transplantation rejection, pulmonary infections, bronchitis,  
 CC and/or cancer. AAF18434 to AAF21543 represent human polynucleotide  
 CC fragments and antisense oligonucleotides used in the exemplification of  
 CC the present invention.  
 XX  
 SQ Sequence 143068 BP; 41194 A; 30122 C; 32403 G; 39349 T; 0 other;

Query Match 56.0%;  
 Best Local Similarity 99.8%;  
 Pred. No. 0;  
 Matches 1252; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 979 AGCCATTTCATAGCTCTTGGCTTGTAGGATGCTCCACTCCAAACCTGTTGCGA 1038  
 Db 48253 AGCCATTTCATAGCTCTTGGCTTGTAGGATGCTCCACTCCAAACCTGTTGCGA 48312  
 Qy 1039 GGTCAGGAGTGAACCGGAAAGAATGACTACACAGGACTCTCGATGG 1098

Db	48313	GGTCAGAGTGTGAGACCAGGAAAGATGTGAAAGTGACTACACAGGACTCCTCGATGGT	48372	Qy	2179	TTTACAACATGATTGGAAAATAATCAATGCTATAACTATGTTGATAAAAG	2232
Qy	1099	CGTGAAAAGAAAAGTCAATTGGAGCCCTGTAAGCAGTCCTAGGAAAGAAAGGA	1158	Db	49453	TTTACAACATGATTGGAAAATAATCAATGCTATAACTATGTTGATAAAAG	49506
Db	48373	CGTGGAAAAGAAAAGTCAATTGGAGCCCTGTAAGCAGTCCTAGGAAAGAAAGGA	48432				
Qy	1159	GCTAGAGACAGAAATGACAGATCTGGTGGAAATCACACGCTGTTGAGTTTACAGATG	12218		RESULT 4		
Db	48433	GCCTAGAGACAGAAATGACAGATCTGGTGGAAATCACACGCTGTTGAGTTTACAGATG	48492		AAA34983		
Qy	1219	TGTGATTCACTAGTGTGAATTCTGGTGTCTACGGTACAGGAGGCTGAGAGAG	1278	ID	143068	BP.	
Db	48493	TGTGATTCACTAGTGTGAATTCTGGTGTCTACGGTACAGGAGGCTGAGAGAG	48552		AAA34983	standard; DNA;	
Qy	1279	AGACTCCAGCCTGGTTGAAACAGTATTCCAACTACCTTCCAGTTCCTCAATTG	1338	XX	28-JUL-2000	(first entry)	
Db	48553	AGACTCCAGCCTGGTTGAAACAGTATTCCAACTACCTTCCAGTTCCTCAATTG	48612	AC			
Qy	1339	AATACAGGCAATAGAGTCAGACTTTTAATAGTAAATAAAATTAAAGCTGAAAC	1398	XX			
Db	48613	AATACAGGCAATAGAGTCAGACTTTTAATAGTAAATAAAATTAAAGCTGAAAC	48672	DT			
Qy	1399	TGCAACTGTAAATGTTGAAAGACTGTGACTCTATCATCTCAAAGTGAAAT	1458	XX			
Db	48673	TGCAACTGTAAATGTTGAAAGACTGTACTATCATCTCAAAGTGAAAT	48732	XX			
Qy	1439	CGTGTATTAGTCAGAGATAATTCTAGCTTGTAGCTTAAAGTTGAGCTGGTAT	1518	OS			
Db	48733	CGTGTATTAGTCAGAGATAATTCTAGCTTGTAGCTTAAAGTTGAGCTGGTAT	48792	XX			
Qy	1519	GTTGGGAGACTGCTGACTCAACCCAATAGTGTGATTGGCAGGACTGGAGTGTG	1578	PN			
Db	48793	GTTGGGAGACTGCTGACTCAACCCAATAGTGTGATTGGCAGGACTGGAGTGTG	48852	XX			
Qy	1579	ATCTGTGGCACATTGGCTATGTCAGTCAGCAACCAATAAGTGTGTTGATG	1638	PD			
Db	48853	ATCTGTGGCACATTGGCTATGTCAGTCAGCAACCAATAAGTGTGTTGATG	48912	XX			
Qy	1639	CAGTATACTCCATCGTGTCACTCAGCTGGATCCTGCACTTCTCAAGGTGTCGCCA	1698	XX			
Db	48913	CAGTATACTCCATCGTGTCACTCAGCTGGATCCTGCACTTCTCAAGGTGTCGCCA	48972	XX			
Qy	1699	AAACGCTTGTGTGTGTATATTAGTACTATGTCAGCTGGTTTAATCACATTGAGT	1758	PT			
Db	48973	AAACGCTTGTGTGTGTATATTAGTACTATGTCAGCTGGTTTAATCACATTGAGT	49032	PT			
Qy	1759	GTTCAGTGTCCGAGTGTCCCTTAATGTCATATTGTCCTCAATTGCGACTGGAA	1818	XX			
Db	49033	GTTCAGTGTCCGAGTGTCCCTTAATGTCATATTGTCCTCAATTGCGACTGGAA	49092	CC			
Qy	1819	CTCCATAATCAAATGGCTCTAACTCAACCTTTAACCTATGGTAAAGAAATGGAA	1878	CC			
Db	49093	CTCCATAATCAAATGGCTCTAACTCAACCTATGGTAAAGAAATGGAA	49152	CC			
Qy	1879	GTGGAGAGTCTCCCTGAAGTAAGCAAGACTTCTCTTGTGCAAGTTAAAGATG	1938	CC			
Db	49153	GTGGAGAGTCTCCCTGAAGTAAGCAAGACTTCTCTTGTGCAAGTTAAAGATG	49212	CC			
Qy	1939	TCCTATGTCGCCAGTGTCTGATCTGATGTCAGTCAGCTGGCTCTCTAG	1998	CC			
Db	49213	TCCTATGTCGCCAGTGTCTGATCTGATGTCAGTCAGCTGGCTCTCTAG	49272	CC			
Qy	1999	ACCGCAACCTGGAACTAGCTCCAGCTTGTGCAAGCTTCACTTCAGGCCACAT	2058	CC			
Db	49273	ACCGCAACCTGGAACTAGCTCCAGCTTGTGCAAGCTTCACTTCAGGCCACAT	49332	CC			
Qy	2059	GGCTAAAGAAAGCTTCAGAAGAAAGCTGGCAAGCTTCACTTCAGGCCACAT	2118	CC			
Db	49333	GGCTAAAGAAAGCTTCAGAAGAAAGCTGGCAAGCTTCACTTCAGGCCACAT	49392	CC			
Qy	2119	GTATGATCCTAAATGAATGTCATAAAATGTTAAGTGTGATGAAATGTAATGTT	2178	CC			
Db	49393	GTATGATCCTAAATGAATGTCATAAAATGTTAAGTGTGATGAAATGTAATGTT	49452	CC			

XX	SQ Sequence 143068 BP; 41194 A; 30122 C; 32402 G; 39350 T; 0 other;	Db 49213 TCTCTATGTTGCCCAAGATGTTCTGATCTGATGAAAGAACACTGGCTTCCTAGA 49272
Query Match	56 0%;	Qy 1999 ACCAGGCACCTTGGAACTTAAAGCTCCAACTGCTTCAACTTCAGGCCACAT 2058
Best Local Similarity	99.8%;	Pred. No. 0;
Matches 1252;	Conservative 0;	Db 49273 ACCAGGCACCTTGGAACTTAAAGCTCCAACTGCTTCAACTTCAGGCCACAT 49332
Qy 979 AGCCCTTTACATAGGCTCTGGCTGTAGATTGCCACTCCAAAACCAAGTGTGGA 1038	Qy 2059 GGCTAAAGAAGGTTTCAGAAAGAAGTGGGACACAGCAGACTTCACCTTCATATT 2118	
Db 48253 AGCCCTTTACATAGCTTGGCTGTAGATTGCCACTCCAAAACCAAGTGTGGA 48312	Db 49333 GGTTAAAGGGTTTCAGAAAGAAGTGGGACACAGCAGACTTCACCTTCATATT 49392	
Qy 1039 GTCAGGGTGTGAGACAGAAAGATGTGAAGTGTACACAGGACTCCTGATG 1098	Qy 2119 GATGATGCTTAATGATGCAAAATGTTAAGTGTGATGTGATGTTAAATGTT 2178	
Db 48313 GTCAGGGTGTGAGACAGAAAGATGTGAAGTGTACACAGGACTCCTGATG 48372	Db 49393 GATGATGCTTAATGATGCAAAATGTTAAGTGTGATGTGATGTTAAATGTT 49452	
Qy 1099 CGTGAAGAAGGAAAGTGTCAAGGACAAAGAAGGA 1158	Qy 2179 TTAACTACTATGTTGGAAATAATTAATGTTGATCTGATAAAAG 2232	
Db 48373 CGTGAAGAAGGAAAGTGTCAAGGACGGCTGAAGCAGAAAGTGTACACAGGACTCCTGATG 4832	Db 49453 TTAACTACTATGTTGGAAATAATTAATGTTGATCTGATAAAAG 49506	
Qy 1159 GCTAGACAGAAATGACAGATCTCTGTTGGAAATCACAGTCAGCTGCTTCACAGAG 1218	RESULT 5	
Db 48433 GCTAGACAGAAATGACAGATCTCTGTTGGAAATCACAGTCAGCTGCTTCACAGAG 48492	AAA35150 standard; DNA; 143068 BP.	
Qy 1219 TGTTGATTCAAGTGTGAACTCTTGTTGTTACGTTACAGGCCAGGGTGTGAGGAG 1278	XX	
Db 48493 TGTTGATTCAAGTGTGAACTCTTGTTGTTACGTTACAGGCCAGGGTGTGAGGAG 48552	AC AAA35150;	
Qy 1279 AGACTCCAGCTGGTTGCAACAGTATTCCAAACTTCCAGTCTGTTTCATTTTG 1338	XX	
Db 48553 AGACTCCAGCTGGTTGCAAAACAGTATTCCAAACTTCCAGTCTGTTTCATTTTG 48612	DE Human adenosine receptor related polynucleotide 2nd SEQ ID NO:24.	
Qy 1339 AATACAGGCTATGAGTTGACTCTTTAAATAGTAAATAAAATTAAGCTGAAAC 1398	XX	
Db 48613 AATACAGGCTATGAGTTGACTCTTTAAATAGTAAATAAAATTAAGCTGAAAC 48672	KW Human adenosine receptor; low adenosine antisense oligonucleotide;	
Qy 1399 TGCAACTGTAAATGTTGAAAGTGTAGTTGAGTTGCTATCATGTCAAACGTGAAAT 1458	KW phosphorothioate; impaired respiration; inflammation; allergy;	
Db 48673 TGCAACTGTAAATGTTGAAAGTGTAGTTGAGTTGCTATCATGTCAAACGTGAAAT 48732	KW allergic disease; bronchoconstriction; inhibitor; antiinflammatory;	
Qy 1459 GCTGTTATTAGTCACAGAGATAATTCTGAGCTTGGAGTTGAGCAGTTGGTA 1518	KW antiallergic; antiasthmatic; cytostatic; analgesic; impaired airway;	
Db 48733 GCTGTTATTAGTCACAGAGATAATTCTGAGCTTGGAGTTGAGCAGTTGGTA 48792	KW lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;	
Qy 1519 GTTTGGGAGACTGGCAGTCAACCCCATAGTGTGTTGAGGTGGAGTTGGTG 1578	KW respiratory distress syndrome; pain; cystic fibrosis; emphysema;	
Db 48793 GTTTGGGAGACTGGCAGTCAACCCCATAGTGTGTTGAGGTGGAGTTGGTG 48852	KW pulmonary hypertension; chronic obstructive pulmonary disease; COPD;	
Qy 1579 ATCTGTGGGACATTAAGCCATTAGCCATTAGTGTGTTGAGGTGGAGTTGGTG 1638	KW cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.	
Db 48853 ATCTGTGGGACATTAAGCCATTAGCCATTAGTGTGTTGAGGTGGAGTTGGTG 48912	XX	
Qy 1639 CAGTATACGGTCCATCGTGTCACTCGTGGGATCTCCATTCTCAGGTTGGCCA 1698	OS Homo sapiens.	
Db 48913 CAGTATACGGTCCATCGTGTCACTCGTGGGATCTCCATTCTCAGGTTGGCCA 48972	XX	
Qy 1699 AAAGCCTTTGTGTTGTTGTTGTTGATCATGTCATGTCATGTCATGTCATGTC 1758	PN WO200009535-A2.	
Db 48973 AAAGCCTTTGTGTTGTTGTTGATCATGTCATGTCATGTCATGTCATGTC 49032	XX	
Qy 1759 GTTTCAGTGTGTCAGATGTTGTTGTTGTTGATCATGTCATGTCATGTCATGTC 1818	PD 24-FEB-2000.	
Db 49033 GTTTCAGTGTGTCAGATGTTGTTGTTGATCATGTCATGTCATGTCATGTC 49092	XX	
Qy 1819 CTCTAAATCAAATGGCTTAATCAAACTTTAAACCTTAACTCATGTCATGTCATGTC 1878	PF 03-AUG-1999;	
Db 49093 CTCTAAATCAAATGGCTTAATCAAACTTTAAACCTTAACTCATGTCATGTCATGTC 49152	XX	
Qy 1879 GTGGAGAGCTCCCTGAACTAAGCTTAACTCATGTCATGTCATGTCATGTCATGTC 1938	PR 03-AUG-1998;	
Db 49153 GTGGAGAGCTCCCTGAACTAAGCTTAACTCATGTCATGTCATGTCATGTCATGTC 49212	XX	
Qy 1939 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	PA (UYEC-) UNIV EAST CAROLINA.	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	PI Nyce JW;	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	XX	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	DR WPI; 2000-205971/18.	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	XX	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	PS Disclosure; Page 1106-1138; 1343pp; English.	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	XX	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC The present invention describes a new composition comprising an	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC antisense oligonucleotide (ON) with low adenosine (up to 15%), which	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC targets nucleic acids involved in bronchoconstriction, allergies, and/or	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC inflammation. The ON can have antiinflammatory, antiallergic,	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC antiasthmatic, cytostatic and analgesic activities. The compositions are	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC used for the treatment of diseases associated with inflammation,	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC impaired airways, including lung disease and diseases whose secondary	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC effects afflict the lungs of a subject. They can be used for treating	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC e.g. ischaemic conditions, pulmonary vasoconstriction, allergies,	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC fibrosis, impaired respiration, respiratory distress syndrome, pain, cystic	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC fibrosis, pulmonary hypertension, emphysema, chronic obstructive	
Qy 1999 TTCTTATGTTGCCAGATGTTCTGATCTGATGTCAGCAAGAACACTGGCTTCCTAGA 1998	CC pulmonary disease (COPD), and cancers such as leukaemias, lymphomas,	

carcinomas, and cancers which may metastasise to the lungs, including breast and prostate cancer. The reduction of the adenosine content of the ONS reduces side effects. The A-containing ONS break down with the release of deoxyadenosine which activates adenosine receptors causing bronchoconstriction and inflammation. AAA3313 to AAA3512 represent the nucleotide sequences given in the sequence listing from the present invention, which correspond to SEQ ID NO:1 to 2815, and then the last 185 sequences are also called SEQ ID NO:1 to 185, but the sequences differ from the previously named sequences. SEQ ID NO:11 to 1680 (AAA3233 to AAA33992) are specifically claimed ONS from the present invention. N.B. Sequences given in the disclosure of the present invention do not match up with their corresponding SEQ ID NO: sequences given in the sequence listing.

Db	4 9093	GTTTCACTGCTTCGAATGTCCTTGTGCTATATGTTCCCTATTGTCAGTGGAA 4 9092
Qy	1819	CTCCTAATCAAATTGGCTCTPAATCAAAGCTTTAACCCATTGCTTAAGATGGAG 1878
Db	4 9093	CTCCTAATCAAATTGGCTCTPAATCAAAGCTTTAACCCATTGCTTAAGATGGAG 4 9152
Qy	1879	GTGGAGAAGCTCCGTAGAGTAAGCAAGACATTCTCTTACTGAGCCAAAGTTAAGATG 1938
Db	4 9153	GTGGAGAAGCTCCGTAGAGTAAGCAAGCTTTAACCCATTGCTTAAGATGGAG 4 9212
Qy	1939	TCTCTATGTTGCCCAAGTGTGTTCTGATCTGTCTGTGCAACCAAGAAACATGGGTTCTAGA 1998
Db	4 9213	TCTCTATGTTGCCCAAGTGTGTTCTGATCTGTCTGTGCAACCAAGAAACATGGGTTCTAGA 4 9272
Qy	1999	ACCGGCAAACCTGGGAACTAGACCTGGGAACCTGGACTATGGCTCTACTTTCACTTCATATATT 2058
Db	4 9273	ACCGGCAAACCTGGGAACTAGACCTGGGAACCTGGACTATGGCTCTACTTTCACTTCATATATT 4 9332
Qy	2059	GGCTAAAGGGTTTCAAGAAGAACTGGGGCACAGGCCAAACTTCACTTCATATATT 2118
Db	4 9333	GGCTAAAGGGTTTCAAGAAGAACTGGGGCACAGGCCAAACTTCACTTCATATATT 4 9392
Qy	2119	GTATGATCCCTAATGATAAAATGTTAACTGTTAACTATGTTAACTATGTT 2178
Db	4 9393	GTATGATCCCTAATGATAAAATGTTAACTATGTTAACTATGTTAACTATGTT 4 9452
Qy	2179	TTTAAACAACTATGATTGGAAATAATGCTAACTATGTTAACTATGTTAACTATGTT 2232
Db	4 9453	TTTAAACAACTATGATTGGAAATAATGCTAACTATGTTAACTATGTTAACTATGTT 4 9506
RESULT <sup>6</sup>		
ID	ABL68124	standard; DNA; 143068 BP.
XX	AC	ABL68124;
XX	DT	15-MAY-2002 (first entry)
XX	DE	Ovary cancer related gene sequence SEQ ID NO:6461.
XX	KW	Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
XX	KW	stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
XX	KW	cytostatic; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma
XX	KW	gene; ds.
OS		Homo sapiens.
XX		WO200194629-A2.
XX		13-DEC-2001.
XX		30-MAY-2001; 2001WO-US10838.
XX	PR	05-JUN-2000; 2000US-209473P.
PR		05-JUN-2000; 2000US-209531P.
PR		18-SEP-2000; 2000US-233133P.
PR		18-SEP-2000; 2000US-233617P.
PR		20-SEP-2000; 2000US-24009P.
PR		20-SEP-2000; 2000US-234034P.
PR		20-SEP-2000; 2000US-234052P.
PR		22-SEP-2000; 2000US-234509P.
PR		22-SEP-2000; 2000US-234567P.
PR		25-SEP-2000; 2000US-234923P.
PR		25-SEP-2000; 2000US-234924P.
PR		25-SEP-2000; 2000US-235077P.
PR		25-SEP-2000; 2000US-235082P.
PR		25-SEP-2000; 2000US-235134P.
PR		25-SEP-2000; 2000US-235280P.
PR		26-SEP-2000; 2000US-235637P.
PR		26-SEP-2000; 2000US-235638P.
PR		27-SEP-2000; 2000US-235711P.

PR	27-SEP-2000;	20000US-235720P.	QY	1159	GCCTAGAGAGAAATGACAGATCTGGCTTGGAAATCACAGCTGGCTCACAGATG 1.218
PR	27-SEP-2000;	20000US-235840P.	Db	48433	GCCAGAGAGAAATGACAGATCTGGCTTGGAAATCACAGCTGGCTCACAGATG 4.8492
PR	28-SEP-2000;	20000US-235865P.	QY	1219	TGTGATTCACTGGAATCTGGCTAGTGTACGGAGGCTGAGAGGAG 1278
PR	28-SEP-2000;	20000US-236028P.	Db	48493	TGTGATTCACTGGAATCTGGCTAGTGTACGGAGGCTGAGAGGAG 4.8552
PR	28-SEP-2000;	20000US-236033P.	QY	1279	AGACTCCACCTGGTTGAAACAGTATTCCAAACTACCTCCAGTCTCATTCTTG 13.38
PR	28-SEP-2000;	20000US-236034P.	Db	48553	AGACTCCACCTGGTTGAAACAGTATTCCAAACTACCTCCAGTCTCATTCTTG 4.8612
PR	28-SEP-2000;	20000US-236109P.	QY	1339	AATACAGGGATAGGTTGAGCTTTAACAGTTAAATGAAATTAAGCTGAAAC 1.398
PR	28-SEP-2000;	20000US-236111P.	Db	48613	AATACAGGGATAGGTTGAGCTTTAACAGTTAAATGAAATTAAGCTGAAAC 4.8672
PR	29-SEP-2000;	20000US-236812P.	QY	1399	TGCAACTTGTAATGTGGTAAGGTTAGTTGGTTGTTGATGTCAACTGAAAT 14.58
PR	02-OCT-2000;	20000US-237171P.	Db	48673	TGCAACTTGTAATGTGGTAAGGTTAGTTGGTTGTTGATGTCAACTGAAAT 4.8732
PR	02-OCT-2000;	20000US-237173P.	QY	1459	GCTGTTAGTCAAGAGATAATTCTAGGTTGGCTTAAAGATTGAGCTTGGTAT 15.18
PR	02-OCT-2000;	20000US-237178P.	Db	48733	GCTGTTAGTCAAGAGATAATTCTAGGTTGGCTTAAAGATTGAGCTTGGTAT 4.8792
PR	02-OCT-2000;	20000US-23728P.	QY	1519	GTTTGGAGACTGCTGAGTCAACCAATAGTTGTTGATGGAGAGTGGAGTGGTGG 15.78
PR	02-OCT-2000;	20000US-237595P.	Db	48793	GTTTGGAGACTGCTGAGTCAACCAATAGTTGTTGATGGAGAGTGGAGTGGTGG 4.8852
PR	03-OCT-2000;	20000US-237716P.	QY	1579	ATCTGTGGCACATAGCCATGCTGATGGCTATGCTAGTGTGTTGAACTGATCA 16.48
PR	03-OCT-2000;	20000US-237725P.	Db	48853	ATCTGTGGCACATAGCCATGCTGATGGCTATGCTAGTGTGTTGAACTGATCA 4.8912
PR	03-OCT-2000;	20000US-237798P.	QY	1639	CAGTAAACGCTCCATGGCTCATCTGCTAGTGTGATGGCTCATCTGCTGGCC 16.98
PR	03-OCT-2000;	20000US-237804P.	Db	48913	CAGTAAACGCTCCATGGCTCATCTGCTAGTGTGATGGCTCATCTGCTGGCC 4.8972
PR	03-OCT-2000;	20000US-237806P.	QY	1699	AAAGCCTTTGTTGTTGTTGTTGATCATATGAACTCATCTGCTGAGT 17.58
PR	01-NOV-2000;	20000US-237908P.	Db	48973	AAAGCCTTTGTTGTTGTTGTTGATCATATGAACTCATCTGCTGAGT 4.9032
PR	01-NOV-2000;	20000US-2444667P.	QY	1759	GTTCAGTGCTGGATGCTCTGATGCTCATATGTTGCCAGTGSSAA 18.18
XX			Db	49033	GTTCAGTGCTGGATGCTCTGATGCTCATATGTTGCCAGTGSSAA 4.9092
PA	(AVAL- AVALON PHARM.				
XX	Young PE, Augustus M,	Carter KC,	Endress G,	Horrigan S;	
PI	Soppet DR,	Weaver Z;			
XX	WPT;	2002-188264/24.			
PT	Screening for anti-neoplastic agent involves exposing cells to a chemical agent to be tested for anti-neoplastic activity, and determining a change in expression of a signature gene set -				
XX	PT	Claim 1; SEQ ID 6461; 44pp; English.			
PS	The present invention describes a method (M1) for screening for an anti-neoplastic agent. The method involves exposing cells to a chemical agent to be tested for anti-neoplastic activity, determining a change in expression of at least one gene (I) of a signature gene set, where (I) comprises a sequence (S) selected from 8471 sequences (given in ABL6164 to AB70110), or is at least 95% identical to (S), where a change in expression is indicative of anti-neoplastic activity. M1 has cytostatic activity and can be used in gene therapy. M1 can be used for screening an anti-neoplastic agent, and can be used for producing a product which is the data collected with respect to the anti-neoplastic agent as a result of M1, and the data is sufficient to convey the chemical structure and/or properties of the agent. M1 can be used in the treatment of cancer such as, colon, breast, stomach, lung, thyroid, oesophageal, ovarian, kidney, prostate or pancreatic cancer, adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer, carcinoma, papillary carcinoma and Wilms' tumour.				
CC	Sequence 143068 BP; 41194 A; 30122 C; 32403 G; 39349 T; 0 other;				
CC	Query Match 56.0%; Score 1250.8; DB 24; Length 143068;				
CC	Best Local Similarity 99.8%; Pred. No. 0; Mismatches 0; Indels 2; Gaps 0;				
CC	Matches 1252; Conservative 0; Mismatches 0;				
CC	QY 979 AGCCCTTTACATAGCTCTGGCTGTAGGATTGCCCACTCCAAAACAGTGTGG 1038				
CC	Db 48253 AGCTTTTACATAGCTCTGGCTGTAGGATTGCCCACTCCAAAACAGTGTGG 4.8312				
CC	QY 1039 GGTCAGGAGTGAGCAGGAAATGGAAGTGTGACTACAGGACCTCGTGTG 1098				
CC	Db 48313 GGTCAGGAGTGAGCAGGAAATGGAAGTGTGACTACAGGACCTCGTGTG 4.8372				
CC	QY 1099 CTCGAAAGGAAAGTCAATTGGAGGCCCTGAGGAAACAGTGTGGCTCTAGA 1158				
CC	Db 48373 CTCGAAAGGAAAGTCAATTGGAGGCCCTGAGGCAAGTGTGGCTCTAGA 4.8432				
CC	QY 2179 TTAAACACTAGTATTTGGAAATTAAATCAATGTTAACTAATGTTGATAAAAG 2232				
CC	Db 49453 TTAAACAACTATGATTGGAAATAATCAATGTTAACTAATGTTGATAAAAG 4.9506				

RESULT 7	
AAA35151	AAA35151 standard; DNA; 149412 BP.
AAA35151;	
XX	28-JUL-2000 (first entry)
XX	Human adenosine receptor related polynucleotide 2nd SEQ ID NO:25.
XX	
XX	Human; adenosine receptor; low adenosine antisense oligonucleotide;
XX	phosphorothioate; impaired respiration; inflammation; allergy;
XX	allergic disease; bronchoconstriction; inhibitor; antiinflammatory;
XX	antiallergic; antiasthmatic; cytostatic; analgesic; impaired airway;
XX	lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;
XX	respiratory distress syndrome; pain; cystic fibrosis; emphysema;
XX	pulmonary hypertension; chronic obstructive pulmonary disease; COPD;
XX	cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.
XX	
XX	Homosapiens.
XX	WO200009525-A2.
XX	24 FEB-2000.
XX	03-AUG-1999; 99WO-US17712.
XX	03-AUG-1998; 98US-0095212.
XX	(UYEC-) UNIV EAST CAROLINA.
XX	NYCE JW;
XX	WPI; 2000-205971/18.
XX	New antisense oligonucleotides useful for treating e.g. pulmonary vasoconstriction, inflammation, allergies, asthma, hypertension, bronchitis, emphysema, respiratory distress syndrome, ischæmia or cancers.
XX	Disclosure; Page 1138-1171; 1343PP; English.
XX	
XX	The present invention describes a new composition comprising an antisense oligonucleotide (ON) with low adenosine (up to 15%), which targets nucleic acids involved in bronchoconstriction, allergies, and/or inflammation. The ON can have antiinflammatory, antiallergic, antasthmatic, cytostatic and analgesic activities. The compositions are useful for the treatment of diseases associated with inflammation, impaired airways, including lung disease and diseases whose secondary effects afflict the lungs of a subject. They can be used for treating e.g. ischæmic conditions, pulmonary vasoconstriction, allergies, asthma, impaired respiration, respiratory distress syndrome, pain, cyst fibrosis, pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), and cancers such as leukaemias, lymphomas, carcinomas, and cancers which may metastasise to the lungs, including breast and prostate cancer. The reduction of the denosine content of the ONs reduces side effects. The A-containing ONs break down with the release of deoxyadenosine which activates adenosine receptors causing bronchoconstriction and inflammation. AA32213 to AA35132 represent the nucleotide sequences given in the sequence listing from the present invention, which correspond to SEQ ID NO:1 to 2815, and then the last 185 sequences are also called SEQ ID NO:1 to 185, but the sequences differ from the previously named sequences. SEQ ID NO:11 to 1680 (AA32323 to AA33992) are specifically claimed ONs from the present invention. N.B. Sequences given in the disclosure of the present invention do not match up with their corresponding SEQ ID NO: sequence given in the sequence listing.
XX	Sequence 149412 BP; 43049 A; 31388 C; 33852 G; 41123 T; 0 other;
XX	Sequence Match 56.0%; Score 1250.8; DB 21; Length 149412;
XX	Best Local Similarity 99.8%; Bred. No. 0;

transmitters, defensins, growth factors, vasoactive peptides and receptors, binding proteins and malignancy associated proteins. The antisense oligonucleotides may be used in this way to treat disorders including respiratory obstruction (especially pulmonary obstruction and/or bronchoconstriction) and/or lung inflammation, allergies) and/or surfactant hypoproduction which are associated with a disease or condition selected from pulmonary vasoconstriction, inflammation, allergies, asthma, impaired respiration, respiratory distress syndrome (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), pulmonary transplantation rejection, pulmonary infections, bronchitis, and/or cancer. AAF1834 to AAF21543 represent human polynucleotide fragments and antisense oligonucleotides used in the exemplification of the present invention.

QY	1759	GTTTCAGTGGCTCCGAGATGCCCTTGTATGCTCATATTGTCAGTGGAA	1818
Db	55377	GTTTCAGTGGCTCCGAGATGCCCTTGTATGCTCATATTGTCAGTGGAA	5543
QY	1819	CTCTTAATCACAATTGGCTCTATCAAAAGCTTAAACCCATTGTTAAAGAATGGAG	1878
Db	55437	CTCTTAATCACAATTGGCTCTATCAAAAGCTTAAACCCATTGTTAAAGAATGGAG	5549
QY	1879	GTGGAAAGTCCCTGAAGTAAGAAGACTTTCCTCTAGTCAGCCAAAGTTAAAGAATG	1938
Db	55497	GTGGAAAGTCCCTGAAGTAAGAAGACTTTCCTCTAGTCAGCCAAAGTTAAAGAATG	5555
QY	1939	TTCTATGTTGCCCAAGTGTGTTCTGATGTCAGGAAACACTGGGCTCTAGA	1998
Db	55557	TTCTATGTTGCCCAAGTGTGTTCTGATGTCAGGAAACACTGGGCTCTAGA	5561
QY	1999	ACCAAGCAACTTGGGAACTAGACTCCAAAGCTGGAACTTGTCTTACCTTCAGGCCACAT	2056
Db	55617	ACCAAGCAACTTGGGAACTAGACTCCAAAGCTGGAACTTGTCTTACCTTCAGGCCACAT	5567
QY	2059	GGCTAAAGAAGTTGAAAGAAGTGGGACAGAGCAGAACTTTCACCTTCATATT	2118
Db	55677	GGCTAAAGAAGTTGAAAGAAGTGGGACAGAGCAGAACTTTCACCTTCATATT	5577
QY	2119	GTATGATCCTTAATGAAATGCAAAATGTTAAGTGTGGCATGAAATCTAAATCTGT	2177
Db	55737	GTATGATCCTTAATGAAATGCAAAATGTTAAGTGTGGCATGAAATCTGT	5577
QY	2179	TTAACAACTATGATTGGAAATAATCAATGCTATAACTATGTTGATAAAAG	2232
Db	55797	TTAACAACTATGATTGGAAATAATCAATGCTATAACTATGTTGATAAAAG	55850

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T      /*tag= a
X      W09519435 - A.
X      20 - JUL - 1995.
X      11 - JAN - 1995; 95WO-US00476.
X      13 - JAN - 1994; 94US-0182962.
X      (REGC ) UNIV CALIFORNIA.
X      Charo I, Coughlin S;
X      WPI; 1995-263866/34 .
X      P-PSDB; AAR79166.
X      DNA encoding monocyte chemo-attractant protein-1 receptor - used par
X      for identifying antagonists and for treating diseases characterised by
X      monocytic infiltrates
X
X      Disclosure; Fig 2; 84PP; English.
X      To identify and clone new members of the chemokine receptor gene

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721 ACCCTGCTTCGTTGCTGAAACGAGAAAGGGCATAGGCAGTGTGAGTCATCTTACCC 780  
 762 ACCCTGCTTCGTTGCTGAAACGAGAAAGGGCATAGGCAGTGTGAGTCATCTTACCC 821  
 781 ATCATGATTGTTACTTCTCTGACTCCCTATACATTGTCATTCCTGAAACCC 840  
 822 ATCATGATTGTTACTTCTCTGACTCCCTATACATGTCATTCCTGAAACCC 881  
 841 TTCCAGGAATCTCGGCTGAGTAACCTGTGAAAGCACCAGTCAACTGGACCAAGCCACGG 900  
 882 TTCCAGGAATCTCGGCTGAGTAACCTGTGAAAGCACCAGTCAACTGGACCAAGCCACGG 941  
 901 CAGGTGACAGAGACTCTGGGTAGACTCAGTGTGCAATCCCATATCTATGCTCTTC 960  
 942 CAGGTGACAGAGACTCTGGGTAGACTCAGTGTGCAATCCCATATCTATGCTCTTC 1001  
 961 GTTGGGGAAAGGTCAAGAG 980  
 1002 GTTGGGGAAAGGTCAAGAG 1021

SULT 10  
312140  
AA512140 standard; DNA: 1083 BP.  
AA512140;  
04-DEC-2001 (first entry)  
Human wild-type CCR2-64V polynucleotide.  
Human: CCR2 receptor; CCR2-64V; CCR2-64V; gene therapy; atherosclerosis; single nucleotide polymorphism; hypercholesterolaemia; ds.

Homo sapiens.  
WO200162796-A1.  
30-AUG-2001.  
22-FEB-2001; 2001WO-GB00755.  
22-FEB-2000; 2000GB-0004183.  
(SMIK) SMITHKLINE BECHAM PLC.  
Valdes AM, Groot PHE, Spurr NK,  
WPI; 2001-550086/61.  
P-PSDE; AAU07614.  
Diagnosing atherosclerosis or susceptibility to atherosclerosis in a

subject, by determining a single nucleotide polymorphism in specific codon of a polynucleotide encoding human CCR2 receptor in genome of the subject -

SQ	Sequence 1083 BP; 255 A; 260 C; 247 G; 321 T; 0 other;
Query Match	42.28%; Score 941; DB 22; Length 1083;
Best Local Similarity	100.0%; Pred. No. 5.5e-52;
Matches	941; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy	40 ATGGTGTCCACATCTCGTGTCTCGGTTATCGAAATAACAGAGAGCGGTTGAAAGTC 99
Db	1 ATGGTGTCCACATCTCGTGTCTCGGTTATCGAAATAACAGAGAGCGGTTGAAAGTC 60
Qy	100 ACCACCTTTTGTATTATGATTAAGGTGTCTCGTGTCTCGGTTATCGAAATAATTGACGTGAAACATT 159
Db	61 ACCACCTTTTGTATTATGATTAAGGTGTCTCGGTTATCGAAATAATTGACGTGAAACATT 120
Qy	160 GGGGCCAACCTCTGCCCTACTCGCTGGTGTCACTCTGGTGTCACTCTGGTGTCAAC 219
Db	121 GGGGCCAACCTCTGCCCTACTCGCTGGTGTCACTCTGGTGTCACTCTGGTGTCAAC 180
Qy	220 ATGCTGGTGTCTCATTTATAAAGCTGAAAAAGCTGAAATGGTGTCTTGTGACTGACATTAC 279
Db	181 ATGCTGGTGTCTCATCTTATAAAGCTGAAAAAGCTGAAATGGTGTCACTGACATTAC 240
Qy	280 CTGCTCAACCTGGCATCTGGTGTCTTGTGATCTGGTTTCTTATTACCTCCATTGTGGCCTAC 339
Db	241 CTGCTCAACCTGGCATCTGGTGTCTTGTGATCTGGTTTCTTATTACCTCCATTGTGGCCTAC 300
Qy	340 TCGTGTCAAATGAGTGGTCTTGGGAATATGGCAAAATTATTCACAGGGCTGTAT 399
Db	301 TCGTGTCAAATGAGTGGTCTTGGGAATGGCAAAATTATTCACAGGGCTGTAT 360
Qy	400 CACATGGTTATTTGGGAAATCTCTTCATCATCTCCCTCAAACTGGATAGATACTCTG 459
Db	361 CACATGGTTATTTGGGAAATCTCTTCATCATCTCCCTCAAACTGGATAGATACTCTG 420
Qy	460 GCTATGGTCATGGTGTGCTTTAAACCGAGCGGTACCTTGGGGTGTGACA 519
Db	421 GCTATGGTCATGGTGTGCTTTAAACCGAGCGGTACCTTGGGGTGTGACA 480
Qy	520 AGTGTGATCACCTGGTGTGCTGTTGCTGTCGCCAGGAATCATCTTACTAAA 579
Db	481 AGTGTGATCACCTGGTGTGCTGTTGCTGTCGCCAGGAATCATCTTACTAAA 540
Qy	580 TGGCAAGAAAGAATGCTGTTATGTCGTCGCCCTATTCACAGGAGTTGGAAATAAT 639
Db	541 TGGCAAGAAAGAATGCTGTTATGTCGTCGCCCTATTCACAGGAGTTGGAAATAAT 600
Qy	640 TTCCACACATAATGAGGACATTTGGGTGCGCTGCTCATCTGGTCATCGTCATGGTCATC 699
Db	601 TTCCACACATAATGAGGACATTTGGGTGCGCTGCTCATCTGGTCATCGTCATGGTCATC 660
Qy	700 TGGTACTCTGGGAATCCTGAAACCCCTGCTGTCGTTGCAACAGAGAGAGGATAGG 759
Db	661 TGGTACTCTGGGAATCCTGAAACCCCTGCTGTCGTTGCAACAGAGAGAGGATAGG 720
Qy	760 GCACTGAGGTCACTTCAACATGATGTTACTTTCTGGACTCCCTPATAAC 819
Db	721 GCACTGAGGTCACTTCAACATGATGTTACTTTCTGGACTCCCTPATAAC 780
Qy	820 ATGGTCACTCTCGAAACCTTCAGGAATTCTGGCCTGAGTAACCTGTAAGGCACC 879
Db	781 ATGGTCACTCTCGAAACCTTCAGGAATTCTGGCCTGAGTAACCTGTAAGGCACC 840
Qy	880 AGTCAACTGGACCAAGCCACGAGGTGACAGAGACTCTGGGTGACTCACTGCTGCATC 939
Db	841 AGTCAACTGGACCAAGCCACGAGGTGACAGAGACTCTGGGTGACTCACTGCTGCATC 900
Qy	940 ATCCCCATCATDATGGCTTCGTTGGGAGAAGTTCAAG 980
Db	901 ATCCCCATCATDATGGCTTCGTTGGGAGAAGTTCAAG 941

RESULT 11

ID	AAS12139	standard; DNA; 1083 BP.	Db	181	ATGCTGGTCATCTCATTAATGCAAAAGCTGAAGTCATGACATTAC	240
XX			QY	280	CTGCTCACCTGGCACACTCTGATCTGGCTTCTTATTACTCTCCATTG	339
AC			Db	241	CTGCTCACCTGGCACAATCTGATCTGGCTTCTTATTACTCTCCATTG	300
XX	04-DEC-2001	(first entry)	QY	340	TCTGCTGCAAATGATGGGGCTTGGGATGCAAATTACAGGGCTGAT	399
XX	Human	CCR2-64I polymorphic variant polynucleotide.	Db	301	TCTGCTGAAATGATGGGGCTTGGGATGCAAATTACAGGGCTGAT	360
XX	KW	Human; CCR2 receptor; CCR2-64I; CCR2-64V; gene therapy; atherosclerosis; single nucleotide polymorphism; hypercholesterolemia; ds.	QY	400	CACATCGGTTATTGGGGAAATCTCTCATATCCCTGACAACTGATAC	459
XX	OS	Homo-sapiens	Db	361	CACATCGGTTATTGGGGAAATCTCTCATATCCCTGACAACTGATAC	420
XX	Key		QY	460	GCTATTGGTCATGTTGTTGCTTAAAGGCCGGACGGTACCTTGGG	519
FT	variation	Location/Qualifiers replace(190,G) /*tag= a /standard_name= "Single nucleotide polymorphism"	Db	421	GCTATTGGTCATGTTGCTTAAAGGCCGGACGGTACCTTGGG	480
FT			QY	520	AGTTGATACCTGTTGGGGCTCTGTTGCTCTGTTGAGGAAATCAT	579
XX	WO200162796-A1.		Db	481	AGTTGATACCTGTTGGGGCTCTGCTGTTGCTGCTCATGAGGATC	540
XX	PD	30-AUG-2001.	QY	580	TGCCAGAAACGAGATTCTGTTTAATGTCAGGATGGATAAT	639
XX	PF	22-FEB-2001; 2001WO-GB00755.	Db	601	TTCACACAAATAATGGAAACCTCTGGCTTGGGCTGTCATGGT	660
XX	PR	22-FEB-2000; 2000GB-0004183.	QY	700	TGCTACTCGGAATCTGAAACCTCTGGCTTGGGCTGCAAAAC	759
XX	PA	(SMIK ) SMITHLINE BEECHAM PLC.	Db	661	TGCTACTCGGAATCTGAAACCTCTGGCTTGGGCTGTCATGG	720
XX	PI	Valides AM, Groot PHE, Spurr NK;	QY	760	GCAGTGAGAGTCATPTCACCATCATGATGTTACTTCTCTGG	819
XX	DR	WPI: 2001-550086/61.	Db	721	GCAGTGAGAGTCATPTCACCATCATGATGTTACTTCTCTGG	780
XX	DR	P-PSDB; AAU07613.	QY	820	ATTTGCTTCTCTGAACTTCTGAACTTCTGAACTTCTGAA	879
XX	PT	Diagnosing atherosclerosis or susceptibility to atherosclerosis in a subject, by determining a single nucleotide polymorphism in specific subject, by determining a single nucleotide polymorphism in specific subject of a polynucleotide encoding human CCR2 receptor in genome of the subject.	Db	781	ATTTGCTTCTCTGAACTTCTGAACTTCTGAACTTCTGAA	840
XX	PT	PT	QY	880	AGTCAACTGGACCAASCCACCGAGACTCTGGATGACTCTGG	939
XX	DR	WPI: 2001-550086/61.	Db	841	AGTCAACTGGACCAAGCCACGGACTCTGGATGACTCTGG	900
XX	XX	XX	QY	940	AATCCCCATCATATGCTTGGGGAGAGTTCAAGAG	980
CC	CC	CC	Db	901	AATCCCCATCATATGCTTGGGGAGAGTTCAAGAG	941
CC	CC	CC	QY	100	ATGCTGGCCACATCTCTTCTGGTTATCAGAAATCCAAACGAG	159
CC	CC	CC	Db	61	ACCACTGGTACATCTCTTCTGGTTATCAGAAATCCAAACGAG	120
CC	CC	CC	QY	160	GGGGCCCAACTCTGGTACATCTCTGGTTATCAGAAATCCAA	219
CC	CC	CC	Db	121	GGGCCCAACTCTGGTACATCTCTGGTTATCAGAAATCCAA	180
CC	CC	CC	QY	220	ATGCTGGTGTCTCATCTTAATGCAAAAGCTGAAGTGTCA	279

Claim 3; Page 20; 28pp; English.

The invention relates to diagnosing atherosclerosis (or susceptibility to) in a subject by determining expression or activity of the human CCR2-64I polypeptide (a polymorphic variant form of the human CCR2 receptor) or the CCR2-64V polypeptide (human CCR2 receptor), by screening for a single nucleotide polymorphism in codon 64 of the polynucleotide encoding the CCR2 receptor. This results in production of CCR2-64I, whereby polymorphic variants are associated with a lower incidence of atherosclerosis. The presence or amount of CCR2-64I/V in a sample can also be analysed. The sequences of the invention can be used for predicting the response of a patient, for predicting the disease outcome in a patient, and also for the production of a treatment for hypercholesterolemia. The sequence represents DNA encoding the polymorphic variant polypeptide CCR2-64I.

Sequence 1083 BP; 256 A; 260 C; 246 G; 321 T; 0 other;

Query Match 42.1%; Score 939.4; DB 22; Length 1083;

Best Local Similarity 99.9%; Pred. No. 1.5e-251; Mismatches 0; Indels 0; Gaps 0;

Matches 940; Conservative 0;

QY 40 ATGCTGGCCACATCTCTTCTGGTTATCAGAAATCCAAACGAG

Db 61 ACCACCTGGTACATCTCTGGTTATCAGAAATCCAAACGAG

QY 100 ACCACCTGGTACATCTCTGGTTATCAGAAATCCAAACGAG

Db 61 ACCACCTGGTACATCTCTGGTTATCAGAAATCCAAACGAG

QY 160 GGGCCCAACTCTGGTACATCTCTGGTTATCAGAAATCCAA

Db 121 GGGCCCAACTCTGGTACATCTCTGGTTATCAGAAATCCAA

QY 220 ATGCTGGTGTCTCATCTTAATGCAAAAGCTGAAGTGTCA

RESULT 12

AAT96976 standard; cDNA; 1083 BP.

XX

AAT96976;

XX

27-FEB-1998 (first entry)

XX

Human monocyte chemoattractant protein 1 receptor encoding cDNA.

XX

Human; MCP-1; monocyte chemoattractant protein; receptor; tumour;

KW inflammatory disease; viral; allergy; diabetes; ds.

XX

Homo sapiens.

XX

Location/Qualifiers

1.1083

/\*tag= a

/product= Monocyte\_chemoattractant\_protein\_1\_receptor

XX

JP09238688-A.



QY 40 ATGCTGICACATCTGTTCTGGTTATCAGAAATACCAACGAGGGTGAAGAGTC 99  
 DE XX Human immune system associated gene SEQ ID NO: 308.  
 DE XX Human; immune system disease; cytosine methylation; antiasthmatic;  
 DE KW antiarteriosclerotic; antianæmic; cytostatic; notropic;  
 DE KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;  
 DE KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;  
 DE KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;  
 DE KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
 DE KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease;  
 DE KW gene; ds.  
 DE XX Homo sapiens.  
 DE XX WO200200928-A2.  
 DE XX 03-JAN-2002.  
 DE XX PD 02-JUL-2001; 2001WO-EP07537.  
 DE XX PF 30-JUN-2000; 2000DE-102529.  
 DE XX PR 01-SEP-2000; 2000DE-1043826.  
 DE XX PA (EPIG-) EPIGENOMICS AG.  
 DE XX PI Olek A, Piepenbrock C, Berlin K;  
 DE XX DR WPI; 2002-130909/17.  
 DE XX PT Nucleic acid comprising fragment of chemically modified gene, useful  
 DE PT for diagnosis and treatment of diseases associated with abnormal  
 DE PT cytosine methylation -  
 DE XX Claim 1; SEQ ID NO 308; 32pp + Sequence Listing; German.  
 DE XX The present invention provides a number of human immune system associated  
 DE CC genes which are modified by the methylation of cytosines. The sequences  
 DE CC can be used in the diagnosis and treatment of immune system disorders,  
 DE CC including eye diseases such as retinopathy, neovascular glaucoma and  
 DE CC macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
 DE CC leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
 DE CC rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
 DE CC diseases. The present sequence is a gene of the invention.  
 DE XX Sequence 10528 BP; 2873 A; 86 C; 2164 G; 5405 T; 0 other;  
 DE SQ Query Match 29.9%; Score 668.2.; DB 24; Length 10528;  
 DE Best Local Similarity 80.2%; Pred. No. 2.6e-175;  
 DE Matches 784; Conservative 0; Mismatches 193; Indels 0; Gaps 0;  
 DE Db 481 ATGTTGATCACCTGGAAATCTGGTGTGTTGGCTTCTGTTGGTAA 540  
 DE QY 580 TGCCAGAAAGAAAGATTCTCTTATGTTGTTGGCCCTTATTTCCAGGAGTGAATAT 639  
 DE Db 541 TGCCAGAAAGAAAGATTCTCTTATGTTGTTGGCCCTTATTTCCAGGAGTGAATAT 600  
 DE QY 640 TTCCACACAAATAATGAGAACATTGGGGCTGGCTCCGGCTCATCATGGTCATC 699  
 DE Db 601 TTCCACACAAATAATGAGAACATTGGGGCTGGCTCCGGCTCATCATGGTCATC 660  
 DE QY 700 TGCTACTGGAAACCTGAAACCTGGCTGGGTGCTGGTGTGCTGGCCAGGAATATCTTACAA 759  
 DE Db 661 TGCTACTGGAAACCTGGCTGGTGTGCTGGTGTGCTGGCTGGATGCTAG 720  
 DE QY 760 GCACTGAGAAGTCACTTCAACCATATGATGTTACTCTCTGGACTCCCTATAAC 819  
 DE Db 721 GCAAAGAGAGTCATCTTCAACCATATGATGTTACTCTCTGGACTCCCTATAAC 780  
 DE QY 820 ATGGTCATCTCCGAAACACCTTCAGGAATTCTGGCTGAACTGTGAAGGACC 879  
 DE Db 781 ATGGTCATCTCCGAAACACCTTCAGGAATTCTGGCTGAACTGTGAAGGACC 840  
 DE QY 880 AGTCAACTGAACTGAACTGGCTGGAGACTCTGGTGAAGTGAAGGCTTC 939  
 DE Db 841 AGTCAACTGAACTGGCTGGAGACTCTGGTGAAGTGAAGGCTTC 900  
 DE QY 940 AATCCCATCATCTATGCTCTGGAGAACTGTTGGCTGAACTGTGAAGGAG 980  
 DE Db 901 AATCCCATCATCTATGCTCTGGAGAACTGTTGGCTGAACTGTGAAGG 941  
 DE QY RESULT 14  
 DE ABL32335/C ID ABL32335 standard; DNA; 10528 BP.  
 DE XX TCTGCCTTTCTTATTAATCTCCATGCTCGTGGCCCTACTCTGCTGAAATGTTGGCTCTT 362  
 DE AC ABL32335;  
 DE XX 26-MAR-2002 (first entry)



QY 661 ATTTCGGCTGGCTCCGGCTGGTCATGATGGTCATCTGGCATACTCGGGAAATCCGAAA 720  
Db 9000 ATTTCGGCTGGCTGGTCATGATGGTCATCTGGCATACTCGGGAAATCCGAAA 9059  
QY 721 ACCCTGCTTCGGTGGTCGAACGAGAAAGGCAATGGCCAGTGAGAGTCATCTTCACCC 780  
Db 9060 ATTTCGGCTTCGGTGGTCGAACGAGAAAGGCAATGGCTAGGTAGAGTTTTATT 9119  
QY 781 ATCATGATGGTTACTTTCTTCGGACTCCCTATAAACATTGTCAATTCTGCATTCTGCACACC 840  
Db 9120 ATTATGGTTATTTGGTTATTTGGATTTTAAATTTGGTATTGGATATT 9179  
QY 841 TTCCAGGATTCCTCGCCTCGAGTACTGAAAGCACAGTCAACTGGACCAAGCCACCG 900  
Db 9180 TTTTAGGAAATTTCGGTTGGAAAGTTAGTTAATTGGATAAGTTACG 9239  
QY 901 CAGGTGACAGAGACTCTGGGATGACTCACAGCTGCAATCCCATCATCTATGCTTC 960  
Db 9240 TAGGTGATAGAGATTGGGATGATTATGGATGATTATTTATTTATGTTTC 9299  
QY 961 GTTGGGGAAAGTTCAAGAGCTTT 987  
Db 9300 GTTGGGGAAAGTTAGAGTT 9326

Search completed: June 1, 2003, 16:03:45  
Job time : 1968.44 secs

